

Examiner-Initiated Interview Summary	Application No.	Applicant(s)
	10/737,350	GEORGES ET AL.
	Examiner	Art Unit
	MISOOK YU, Ph.D.	1642

All Participants:

Status of Application: _____

(1) MISOOK YU, Ph.D.

(3) _____

(2) Dr. David Chavous.

(4) _____

Date of Interview: 1/5/2007

Time: _____

Type of Interview:

Telephonic
 Video Conference
 Personal (Copy given to: Applicant Applicant's representative)

Exhibit Shown or Demonstrated: Yes No

If Yes, provide a brief description: *Applicant's supplemental response, attached to this interview summary.*

Part I.

Rejection(s) discussed:

Claims discussed:

claims 10-108

Prior art documents discussed:

none

Part II.

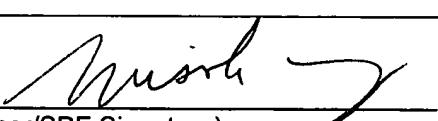
SUBSTANCE OF INTERVIEW DESCRIBING THE GENERAL NATURE OF WHAT WAS DISCUSSED:

The examiner contacted applicant to cancel the withdrawn claims for allowance, and also to amend the specification.

Part III.

It is not necessary for applicant to provide a separate record of the substance of the interview, since the interview directly resulted in the allowance of the application. The examiner will provide a written summary of the substance of the interview in the Notice of Allowability.

It is not necessary for applicant to provide a separate record of the substance of the interview, since the interview did not result in resolution of all issues. A brief summary by the examiner appears in Part II above.


 (Examiner/SPE Signature)

(Applicant/Applicant's Representative Signature – if appropriate)

**FACSIMILE**

Date January 5, 2007**+1 617 526 6523 (I)**
+1 617 526 5000 (F)**david.chavous@wilmerhale.com**

To Examiner Missok Yu
**United States Patent and Trademark
Office**
Fax 571-273-0839
Tel 571-272-0839**cc**

From David A. Chavous, Ph.D. **Pages** 9 (including cover)

Re U.S. Application No. 10/737,350

Our Ref: 112418.149US2/AUR-011US

Dear Examiner Yu:

Attached please find a courtesy copy of the Supplemental Amendment that was faxed today to the official of the USPTO.

Thank you.

Wilmer Cutler Pickering Hale and Dorr LLP, 60 State Street, Boston, Massachusetts 02109
Baltimore Beijing Berlin Boston Brussels London New York Oxford Palo Alto Waltham Washington

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PAGE 1/9 * RCVD AT 1/5/2007 2:39:58 PM [Eastern Standard Time] * SVR:USPTO-EFXRF-5/13 * DNIS:2730839 * CSID:16175265000 * DURATION (mm:ss):01:38

PTO/SB/21 (08-06)

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TRANSMITTAL FORM

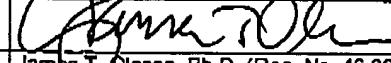
(to be used for all correspondence after initial filing)

TRANSMITTAL FORM (to be used for all correspondence after initial filing)	Application Number	10/737,350-Conf. #6000	
	Filing Date	December 15, 2003	
	First Named Inventor	Elias GEORGES	
	Art Unit	1642	
	Examiner Name	M. Yu	
Total Number of Pages in This Submission	10	Attorney Docket Number	112418.149US2/AUR-011US

ENCLOSURES (Check all that apply)

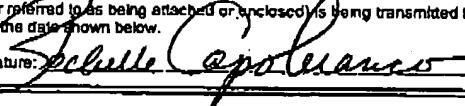
<input type="checkbox"/> Fee Transmittal Form <input type="checkbox"/> Fee Attached <input checked="" type="checkbox"/> Supplemental Response (7 pages) <input type="checkbox"/> After Final <input type="checkbox"/> Affidavits/declaration(s) <input type="checkbox"/> Extension of Time Request <input type="checkbox"/> Express Abandonment Request <input type="checkbox"/> Information Disclosure Statement <input type="checkbox"/> Certified Copy of Priority Document(s) <input type="checkbox"/> Reply to Missing Parts/ Incomplete Application <input type="checkbox"/> Reply to Missing Parts under 37 CFR 1.52 or 1.53	<input type="checkbox"/> Drawing(s) <input type="checkbox"/> Licensing-related Papers <input type="checkbox"/> Petition <input type="checkbox"/> Petition to Convert to a Provisional Application <input type="checkbox"/> Power of Attorney, Revocation Change of Correspondence Address <input type="checkbox"/> Terminal Disclaimer <input type="checkbox"/> Request for Refund <input type="checkbox"/> CD, Number of CD(s) _____ <input type="checkbox"/> Landscape Table on CD	<input type="checkbox"/> After Allowance Communication to TC <input type="checkbox"/> Appeal Communication to Board of Appeals and Interferences <input type="checkbox"/> Appeal Communication to TC (Appeal Notice, Brief, Reply Brief) <input type="checkbox"/> Proprietary Information <input type="checkbox"/> Status Letter <input checked="" type="checkbox"/> Other Enclosure(s) (please identify below): - Fax Transmittal (1 p.) - Certificate of Transmission (1 p.)
Remarks		

SIGNATURE OF APPLICANT, ATTORNEY, OR AGENT

Firm Name	WILMER CUTLER PICKERING HALE AND DORR LLP		
Signature			
Printed name	James T. Olesen, Ph.D. (Reg. No. 46,967), signing on behalf of Ann-Louise Kerner, Ph.D. (Reg. No. 33,523)		
Date	January 5, 2007	Reg. No.	33,523

I hereby certify that this paper (along with any paper referred to as being attached or enclosed) is being transmitted by facsimile to the Patent and Trademark Office, facsimile no. (571) 273-6300, on the date shown below.

Dated: January 5, 2007

Signature: 

(Rochelle Capobianco)

Application No. 10/737,350
 Attorney Docket No. 112418.149US2/AUR-011US
 Supplemental Response Dated 1/5/2007

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant:	Georges <i>et al.</i>	Art Unit:	1642
Serial No.:	10/737,350	Examiner:	Yu, Misook
Filing Date:	December 15, 2003	Customer No.	23483
Title:	HSC70 Directed Diagnostics And Therapeutics For Multidrug Resistant Neoplastic Disease	Conf. No.	6000

CERTIFICATION UNDER 37 CFR § 1.8(a)

I hereby certify that this correspondence is being communicated to the United States Patent and Trademark Office by Facsimile to Fax No. 571-273-8300.

01-05-2007

Date



Rochelle Capobianco

Mail Stop Amendment
 Commissioner for Patents
 P.O. Box 1450
 Alexandria, VA 22313-1450

SUPPLEMENTAL RESPONSE

Dear Sir:

In response to the Examiner interviews of January 4, 2007 and January 5, 2007, and following Applicant's Amendment and Response to the Office Action dated August 1, 2006, Applicants respectfully submit the following Amendments to conform with formalities as requested by the Examiner.

Amendments to the Specification begin on page 2 of this paper.

Amendments to the Claims begin on page 4 of this paper.

Remarks begin on page 6 of this paper.

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Conclusions begin on page 6 of this paper.

AMENDMENTS TO THE SPECIFICATION

Please delete the paragraph on page 19, lines 7-17 of the application, and replace it with the following paragraph.

In particular, this application incorporates the following patent applications by reference in their entirety: U.S.S.N. 60/433,480, filed Dec. 13, 2002 and entitled "Vimentin Detection-Based Methods for Diagnosing and Treating Damaged Cells, Neoplastic Cells and Multidrug Resistance;" U.S.S.N. 60/433,351, filed Dec. 13, 2002 and entitled "Nucleophosmin Detection-Based Methods for Diagnosing and Treating Damaged Cells, Neoplastic Cells and Multidrug Resistance", as well as U.S.S.N. 10/736,889 YY/XXXXXX, filed Dec. 15, 2003 and entitled "Vimentin Directed Diagnostics and Therapeutics for Multidrug Resistant Neoplastic Disease;" and U.S.S.N. 60/438,012, filed Jan. 1, 2003 and entitled "HSC70 Detection-Based Methods for Diagnosing and Treating Damaged Cells, Neoplastic Cells and Multidrug Resistance," as well as U.S.S.N. 10/737,712 YY/XXXXXX, filed Dec. 15, 2003 and entitled "Nucleophosmin Directed Diagnostics and Therapeutics ~~Therapeutics and Diagnostics~~ for Multidrug Resistant Neoplastic Disease."

Please delete the paragraph on page 76, lines 13-27 of the application, and replace it with the following paragraph.

Examples of pathway-responsive promoters useful in the practice of the present invention include synthetic insulin pathway-responsive promoters containing the consensus insulin binding sequence (Jacob, et al. (1995) *J. Biol. Chem.* 270:27773-27779), the cytokine pathway-responsive promoter, the glucocorticoid pathway-responsive promoter (Lange, et al. (1992) *J. Biol. Chem.* 267:15673-80), IL1 and IL6 pathway-responsive promoters (Won K.-A and Baumann H. (1990) *Mol. Cell. Biol.* 10: 3965-3978), T3 pathway-responsive promoters, thyroid hormone pathway-responsive promoters containing the consensus motif, the TPA pathway-responsive promoters (TREs), TGF-beta pathway-responsive promoters (as described in Grotendorst, et al. (1996) *Cell Growth and Differentiation* 7: 469-480). Additionally, natural or synthetic E2F pathway responsive promoters may be used. An example of an E2F pathway responsive promoter is described in Part, et al. (1997) *Nature Medicine* 3:1145-1149) which describes an E2F-1 promoter containing 4 E2F binding sites and is reportedly active in tumor

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cells with rapid cycling. Examples of other pathway-responsive promoters are well known in the art and can be identified in the Database of Transcription Regulatory Regions on Eukaryotic Genomes accessible through the internet at world wide web eimb.rssi.ru/TRRD <http://www.eimb.rssi.ru/TRRD>.

Please delete the paragraph on page 90, lines 17-25 of the application, and replace it with the following paragraph.

Another method for determining antigenicity of a polypeptide subsequence is the algorithm of Hopp and Woods ((1981) Proc. Natl. Acad. Sci. 86: 152-6). There are publicly available web sites for Hopp and Woods algorithm analysis of a user-input polypeptide sequence and convenient graphical output of the resulting analysis (see, e.g., hypertext transfer protocol http://hometown.aol.com/_ht_a/lucatoldo/myhomepage/JaMBW/3/1/7/). Using this algorithm to analyze the full-length human HSC70 sequence shown in Figure 14A, several suitable sequence having a high Hopp and Woods antigenic index of an adequate length for immunogenicity were revealed. These include HSC70 amino acid residues: 240-260 (i.e. HFIAEFKRKHKKDISENKRAY); and 480-500 (i.e., IDANGILNVSAVDKSTGKENK).

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AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (previously presented) A method for detecting multidrug resistance or multidrug resistance potential in a test neoplastic cell, comprising:
 - a) measuring a level of cell surface-expressed HSC70 protein in the test neoplastic cell of a given origin or cell type; and
 - b) comparing the level of cell surface-expressed HSC70 protein in the test neoplastic cell to the level of cell surface-expressed HSC70 in a nonresistant neoplastic cell of the same origin or cell type,
wherein the test neoplastic cell is multidrug resistant or has multidrug resistance potential if the level of cell surface-expressed HSC70 in the test neoplastic cell is greater than the level of cell surface-expressed HSC70 in the nonresistant neoplastic cell of the same given origin or cell type.
2. (previously presented) The method of claim 1, wherein measuring the level of cell surface-expressed HSC70 in the test neoplastic cell comprises isolating a cytoplasmic membrane fraction from the cell and measuring the level of HSC70 in the cytoplasmic membrane fraction.
3. (previously presented) The method of claim 1, wherein measuring the level of cell surface-expressed HSC70 in the test neoplastic cell comprises contacting said cell with an anti-HSC70 antibody and measuring the level of antibody bound to cell surface HSC70.
4. (previously presented) The method of claim 3, wherein measuring the level of antibody bound to cell surface HSC70 is by immunofluorescence emission.
5. (previously presented) The method of claim 3, wherein measuring the level of antibody bound to cell surface HSC70 is by radiolabel.

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6. (previously presented) The method of claim 1, wherein the test neoplastic cell is selected from the group consisting of a promyleocytic leukemia cell, a T lymphoblastoid cell, a breast epithelial cell, and an ovarian cell.
7. (previously presented) The method of claim 1, wherein the nonresistant neoplastic cell is from a drug-sensitive cell line selected from the group consisting of HL60, NB4, CEM, HSB2 Molt4, MCF-7, MDA, SKOV-3, and 2008.
8. (previously presented) The method of claim 1, wherein the test neoplastic cell is selected from the group consisting of a lymphoma cell, a melanoma cell, a sarcoma cell, a leukemia cell, a retinoblastoma cell, a hepatoma cell, a myeloma cell, a glioma cell, a mesothelioma cell, and a carcinoma cell.
9. (previously presented) The method of claim 1, wherein the test neoplastic cell is from a tissue selected from the group consisting of blood, bone marrow, spleen, lymph node, liver, thymus, kidney, brain, skin, gastrointestinal tract, eye, breast, prostate, and ovary.

10.-108. (cancelled).

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REMARKS

1. *History*

Applicants thank the Examiner for the interviews of January 4, 2007 and January 5, 2007. During the interview of January 4, 2007, the Examiner indicated that the claims were in condition for allowance provided that the Applicants cancelled claims 10-108, which had been withdrawn for prosecution at a later date. The Examiner further stated that amendments to the specification were required prior to allowance.

In the interview of January 5, 2007, the Examiner indicated that Applicants should fax a copy of these amendments to the fax number listed on the Office Action dated August 1, 2006.

2. *Claims And Amendments*

Applicants cancel claims 10-108, and reserve the right to prosecute the claims at a later date. Accordingly, no new matter is introduced by these Amendments.

Applicants also have amended the specification per the Examiner's request. In particular, the embedded hyperlinks on pages 76 and 90 have been removed to conform to 37 C.F.R. 1.57(d). Also, patent application serial numbers have been provided on page 19, lines 7-27. Therefore, the specification has been amended to conform to Patent Office policy and per the Examiner's request. Accordingly, Applicants submit that no new matter is introduced by these Amendments.

CONCLUSIONS

Applicants again thank the Examiner for the telephonic interviews of January 4, 2007 and January 5, 2007. In view of the amendments requested during those interviews, Applicants request favorable consideration of the pending claims.

No additional fees are believed to be due in connection with this response. However, please charge any underpayments or credit any overpayments to Deposit Account No. 08-0219.

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If the Examiner believes that any further discussion of this communication would be helpful, please contact the undersigned at the telephone number provided below.

Respectfully submitted,



Ann-Louise Kerner, Ph.D.
Reg. No. 33,523

James T. Olesen, Ph.D.
Signing for Ann-Louise Kerner, Ph.D.
Reg. No. 46,967

January 5, 2007
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